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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/068,471	02/04/2002	Malcolm Lovell Handel	529282000400	7001
1444 7	590 09/02/2004		EXAMINER	
BROWDY AND NEIMARK, P.L.L.C.			FREDMAN, JEFFREY NORMAN	
624 NINTH ST SUITE 300	TREET, NW		ART UNIT	PAPER NUMBER
	N, DC 20001-5303		1637	

DATE MAILED: 09/02/2004

Please find below and/or attached an Office communication concerning this application or proceeding.

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	Application No.	Applicant(s)			
	10/068,471	HANDEL ET AL			
Office Action Summary	Examiner	Art Unit			
	Jeffrey Fredman	1637			
The MAILING DATE of this communication appears on the cover sheet with the correspondence address Period for Reply					
A SHORTENED STATUTORY PERIOD FOR REPLY THE MAILING DATE OF THIS COMMUNICATION.  - Extensions of time may be available under the provisions of 37 CFR 1.13 after SIX (6) MONTHS from the mailing date of this communication.  - If the period for reply specified above is less than thirty (30) days, a reply - If NO period for reply is specified above, the maximum statutory period w - Failure to reply within the set or extended period for reply will, by statute, Any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b).	36(a). In no event, however, may a reply be y within the statutory minimum of thirty (30) vill apply and will expire SIX (6) MONTHS fr , cause the application to become ABANDO	timely filed  days will be considered timely.  om the mailing date of this communication.  NED (35 U.S.C. § 133).			
Status					
<ul> <li>1) Responsive to communication(s) filed on</li> <li>2a) This action is FINAL. 2b) This action is non-final.</li> <li>3) Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under Ex parte Quayle, 1935 C.D. 11, 453 O.G. 213.</li> </ul>					
Disposition of Claims					
4) ☐ Claim(s) 1-18 is/are pending in the application. 4a) Of the above claim(s) is/are withdray 5) ☐ Claim(s) is/are allowed. 6) ☐ Claim(s) is/are rejected. 7) ☐ Claim(s) is/are objected to. 8) ☐ Claim(s) 1-18 are subject to restriction and/or expressions.	vn from consideration.				
Application Papers					
9) The specification is objected to by the Examine 10) The drawing(s) filed on is/are: a) access Applicant may not request that any objection to the of Replacement drawing sheet(s) including the correction of the oath or declaration is objected to by the Examine	epted or b) objected to by the drawing(s) be held in abeyance. So ion is required if the drawing(s) is	See 37 CFR 1.85(a). objected to. See 37 CFR 1.121(d).			
Priority under 35 U.S.C. § 119					
12) ☐ Acknowledgment is made of a claim for foreign a) ☐ All b) ☐ Some * c) ☐ None of:  1. ☐ Certified copies of the priority documents 2. ☐ Certified copies of the priority documents 3. ☐ Copies of the certified copies of the prior application from the International Bureau * See the attached detailed Office action for a list of	s have been received. s have been received in Applic rity documents have been rece u (PCT Rule 17.2(a)).	ation No ived in this National Stage			
Attachmont/c)					
Attachment(s)  1) Notice of References Cited (PTO-892)  2) Notice of Draftsperson's Patent Drawing Review (PTO-948)  3) Information Disclosure Statement(s) (PTO-1449 or PTO/SB/08)	4)  Interview Summa Paper No(s)/Mail 5)  Notice of Informa	ary (PTO-413) Date Il Patent Application (PTO-152)			

Paper No(s)/Mail Date \_\_\_\_\_.

6) Other: \_\_\_\_\_

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#### **DETAILED ACTION**

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#### Election/Restrictions

- 1. Restriction to one of the following inventions is required under 35 U.S.C. 121:
  - I. Claims 1-9, drawn to a DNAzyme, classified in class 435, subclass 18.
  - II. Claims 10-11, 17 and 18, drawn to methods of inhibiting NF-kB, classified in class 514, subclass 44.
  - III. Claims 12-16, drawn to methods of treating disease, classified in class514, subclasses 825, 921.

The inventions are distinct, each from the other because of the following reasons:

- 2. Inventions in Group I and in Groups II and III are related as product and process of use. The inventions can be shown to be distinct if either or both of the following can be shown: (1) the process for using the product as claimed can be practiced with another materially different product or (2) the product as claimed can be used in a materially different process of using that product (MPEP § 806.05(h)). In the instant case, the DNAzyme of Group I can be used in NF-kB inhibition methods as in Group II, in methods of treating disease, in nucleic acid digestion methods, in nucleic acid purification methods or in enzymatic assay methods.
- 3. Inventions in Group II and Group III are unrelated. Inventions are unrelated if it can be shown that they are not disclosed as capable of use together and they have different modes of operation, different functions, or different effects (MPEP § 806.04, MPEP § 808.01). In the instant case the different inventions are unrelated because

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Group II differs in mode of operation, function and effect from Group III, since Group II results in an inhibition of NF-kB while Group III results in a treatment of a disease.

- 4. Because these inventions are distinct for the reasons given above and have acquired a separate status in the art as shown by their different classification, restriction for examination purposes as indicated is proper. Also, these inventions are distinct for the reasons given above and the search required for each Group is distinct, since a search for a disease treatment is a different search than that of an inhibitor and both differ from the search required for a product which may not operate in either way in the prior art yet remain anticipated.
- 5. This application contains claims directed to the following patentably distinct species of the claimed invention:

Two separate species elections are required, which depend in part on the group that is elected. If Group I is elected, then the first species election is the only one required, and applicant is required to elect one target site and the corresponding sequence of one DNAzyme for initial examination under Markush practice. If Group III is elected, Applicant is required to make an additional species election of a disease which is treated by the method.

### Species election 1

The species are the Groups (i)-(xxxv) in claim 4 and SEQ ID NO: 2, SEQ ID Nos: 3-37 in claim 6 and the sequence in claim 7 that lacks a SEQ ID NO (which should be added by amendment).

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Applicant is required under 35 U.S.C. 121 to elect a single disclosed species, which may represent one DNAzyme sequence from one of claims 2, 6 and 7 and the corresponding target site from claim 4, for prosecution on the merits to which the claims shall be restricted if no generic claim is finally held to be allowable. Currently, claims 1-2 and 8-18 are generic.

## **Species election 2**

The species are the diseases listed in claims 13-16.

Applicant is required under 35 U.S.C. 121 to elect a single disclosed species, which may represent one disease such as asthma, or septic shock or leukemia, for prosecution on the merits to which the claims shall be restricted if no generic claim is finally held to be allowable. Currently, claims 12, 17 and 18 are generic.

Applicant is advised that a reply to this requirement must include an identification of the species that is elected consonant with this requirement, and a listing of all claims readable thereon, including any claims subsequently added. An argument that a claim is allowable or that all claims are generic is considered nonresponsive unless accompanied by an election.

Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to additional species which are written in dependent form or otherwise include all the limitations of an allowed generic claim as provided by 37 CFR 1.141. If claims are added after the election, applicant must indicate which are readable upon the elected species. MPEP § 809.02(a).

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Should applicant traverse on the ground that the species are not patentably distinct, applicant should submit evidence or identify such evidence now of record showing the species to be obvious variants or clearly admit on the record that this is the case. In either instance, if the examiner finds one of the inventions unpatentable over the prior art, the evidence or admission may be used in a rejection under 35 U.S.C. 103(a) of the other invention.

- 6. Applicant is advised that the reply to this requirement to be complete must include an election of the invention to be examined even though the requirement be traversed (37 CFR 1.143).
- 7. Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a request under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Jeffrey Fredman whose telephone number is (571)272-0742. The examiner can normally be reached on 6:30-4:00.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Gary Benzion can be reached on (571)272-0782. The fax phone number for the organization where this application or proceeding is assigned is 703-872-9306.

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Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

Jeffrey Fredman Primary Examiner Page 6

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